Application No. 10/527,414

Amd. Dated: November 19, 2009

Reply to Office Action mailed August 25, 2009

Amendment to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1 (currently amended): A medical implant for the controllable delivery of at least one pharmaceutical compound to a localized area within a patient, said implant comprising:

an implantable medical device having a surface and a coating formed on at least a portion of said surface, said coating having at least two polymer layers, two of said at least two polymer layers incorporating at least one releasable pharmaceutical compound, each of said two polymer layers incorporating at least one releasable pharmaceutical compound having at least one physical property affecting the releasability of said releasable pharmaceutical compound that differs from said other layer, wherein said at least one physical property affecting the releasability of said at least one pharmaceutical compound is molecular weight and wherein said at least one releasable pharmaceutical compound is a macrolide antibiotic.

Claim 2 (original): The medical implant of claim 1 wherein said medical device is selected from the group consisting of stents, probes, catheters, micro-particles, pacing leads, vascular grafts, access devices, in-dwelling access ports, valves, plates, barriers, supports, shunts, dises, and joints.

Claim 3 (original): The medical implant of claim 2 wherein said stent is selected from the group consisting of vascular stents, biliary stents, and esophogeal stents.

Claims 4-5 (canceled).

Claim 6 (previously presented): The medical implant of claim 1 wherein said molecular weight range from about 1 kDa to 100,000 kDa.

Claim 7 (previously presented): The medical implant of claim 1 wherein said polymer layers comprise a polymer is selected from the group consisting of poly(caprolactone), poly(lactic acid), poly(glycolic acid), poly(ethylene-vinyl acetate), collagen, heparinized collagen, polyvinyl pyrrolidone, polytetrafluoroethylene, polyethylene glycol, polystyrene,

Application No. 10/527,414 Amd. Dated: November 19, 2009

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acrylates, polyesters, epoxides, silicones, cellulose, and copolymers thereof.

Claim 8 (canceled)

Claim 9 (canceled).

Claim 10 (currently amended): The medical implant of claim 9 1 wherein the

macrolide antibiotic is rapamycin or analogues and derivatives thereof.

Claim 11 (canceled)

Claim 12 (previously presented): A method for making a controllable drug

releasing gradient coating for the surface of a medical device, said method comprising the steps

of:

forming a first polymer layer on said surface of said medical device, said first

polymer layer containing at least one releasably bound pharmaceutical compound and having at

least one physical property affecting the releasability of said at least one pharmaceutical

compound; and

forming at least one additional polymer layer on said first polymer layer, said at

least one additional layer containing at least one releasably bound pharmaceutical compound,

said additional polymer layer differing in said at least one physical property affecting the

releasability of said at least one pharmaceutical compound from said first polymer layer, wherein

said at least one physical property affecting the releasability of said at least one pharmaceutical

compound is molecular weight and wherein the at least one releasably bound pharmaceutical

compound is a macrolide antibiotic.

Claim 13 (original): The method of claim 12 wherein said generally tubular

structure is a stent or a catheter.

Claim 14 (original): The method of claim 13 wherein said stent is self-expanding.

Claim 15 (original): The method of claim 13 wherein said stent is mechanically

Page 3 of 8

Application No. 10/527,414 Amd. Dated: November 19, 2009

Reply to Office Action mailed August 25, 2009

expandable.

Claim 16 (original): The method of claim 13 wherein said stent is bioresorbable.

Claim 17 (canceled).

Claim 18 (previously presented): The method of claim 12 wherein said molecular weights range from about 1 kDa to 100,000 kDa.

Claim 19 (original): The method of claim 12 wherein said polymer layers are selected from the group consisting of poly(caprolactone), poly(lactic acid), poly(glycolic acid), poly(ethylene-vinyl acetate), collagen, heparinized collagen, polyvinyl pyrrolidone, polytetrafluoroethylene, polyethylene glycol, polystyrene, acrylates, polyesters, epoxides,

silicones, cellulose, and copolymers thereof.

Claim 20 (currently amended): The method of claim 12 wherein said at least one anti-restenotic releasably bound pharmaceutical compound is contained within adjacent polymer coatings.

Claim 21 (canceled)

Claim 22 (currently amended): The method of claim 21 20 wherein the macrolide antibiotic is rapamycin or analogues and derivatives thereof.

Claim 23 (currently amended): The method of claim 12 wherein said at least one anti-restenotie releasably bound pharmaceutical compound is coupled to said polymer coating.

Claim 24 (canceled)

Claim 25 (currently amended): The method of claim 24 23 wherein the macrolide antibiotic is rapamycin or analogues and derivatives thereof.